

DETAILED ACTION

Application status

Claims 1-85 are pending in this application.

Priority

The instant application is the 371 national stage entry of PCT/US05/11821, filed on 04/07/2005, which claims benefit of 60/561,110 filed on 04/09/2004 as requested in the declaration.

Election

Applicant's election without traverse of Group III, Claims 38-44 and 55-64, and SEQ ID NOs: 27 (pckA nucleic acid) and 28 (pckA protein), in the response filed on 06/21/11, is acknowledged.

Claims 1-37, 45-54 and 65-85 are withdrawn from further consideration pursuant to 37 CFR 1.142(b) as being drawn to a non-elected invention.

Applicant is reminded that upon the cancellation of claims to a non-elected invention, the inventorship must be amended in compliance with 37 CFR 1.48(b) if one or more of the currently named inventors is no longer an inventor of at least one claim remaining in the application. Any amendment of inventorship must be accompanied by a request under 37 CFR 1.48(b) and by the fee required under 37 CFR 1.17(i).

Information Disclosure Statement

Applicants' filing of information disclosure, filed on 12/03/2007, is acknowledged.
All of the references cited therein have been considered by the Examiner.

Claim Rejections - 35 U.S.C. § 112

The following is a quotation of the second paragraph of 35 U.S.C. § 112:

The specification shall conclude with one or more claims particularly pointing out and distinctly claiming the subject matter which the applicant regards as his invention.

Claims 43, 56 and 61 are rejected under 35 U.S.C. § 112, second paragraph, as being indefinite for failing to particularly point out and distinctly claim the subject matter which applicant regards as the invention.

Claim 43 recites the limitation "at least one chromosomal gene" in Claim 38. There is insufficient antecedent basis for this limitation in the claim. In the interest of advancing prosecution, Claim 43 has been interpreted to depend from Claim 41.

Claim 56 recites the limitation "at least one gene..." in Claim 55. However, the claim is unclear as to whether [1] this is a further comprising step of inactivating one of those genes listed in claim 56, or [2] one of those genes listed in claim 56 is deleted at the same time as the pckA gene. In the interest of advancing prosecution, Claim 56 is interpreted as a further comprising step of inactivating one of those genes listed in claim 56 as suggested by claim 41.

Claim 61 recites the limitation "degU(Hy)32" which is unclear and indefinite. The reason is that the position it is referring to is unclear without a reference sequence, i.e.,

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a SEQ ID NO, to which this position is referring to. Applicants' clarification is suggested.

Claim Rejections - 35 U.S.C. § 102

The following is a quotation of the appropriate paragraphs of 35 U.S.C. § 102 that form the basis for the rejections under this section made in this Office action:

A person shall be entitled to a patent unless –

(a) the invention was known or used by others in this country, or patented or described in a printed publication in this or a foreign country, before the invention thereof by the applicant for a patent.

(e) the invention was described in (1) an application for patent, published under section 122(b), by another filed in the United States before the invention by the applicant for patent or (2) a patent granted on an application for patent by another filed in the United States before the invention by the applicant for patent, except that an international application filed under the treaty defined in section 351(a) shall have the effects for purposes of this subsection of an application filed in the United States only if the international application designated the United States and was published under Article 21(2) of such treaty in the English language.

Claims 38-44 and 55-64 are rejected under 35 U.S.C. § 102(a) as being anticipated by Ferrari et al. (WO/2003/083125 published on 10/09/2003, see IDS).

The instant claims are drawn to [1] a method for enhancing production of at least one protein by a microorganism, comprising the steps: a) providing a microorganism host cell; b) inactivating the *pckA* gene in said host cell to produce an altered strain; and c) growing said altered strain under growth conditions suitable for expression of said protein; and [2] a method for obtaining *Bacillus subtilis* strains having enhanced protease production comprising the steps of: transforming a *B. subtilis* host cell with a DNA construct; allowing homologous recombination of the DNA construct and a homologous region of the *B. subtilis* chromosome wherein *pckA* is deleted from the *B.*

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subtilis chromosome to produce an altered *Bacillus subtilis* strain; and growing the altered *B. subtilis* strain under conditions suitable for the expression of said protease.

It is noted by the Examiner that the term, “homologous proteins” and “heterologous proteins” are defined in page 38 of the specification, and have been interpreted by the Examiner as how they are defined in the specification.

Ferrari et al. teach a method for enhancing expression of a protein of interest from *Bacillus* comprising: a) obtaining an altered *Bacillus* strain capable of producing a protein of interest, wherein said altered *Bacillus* strain has at least one inactivated chromosomal gene selected from the group consisting of *sbo*, *sir*, *ybcO*, *csn*, *spolISA*, *sigB*, *phrC*, *rapA*, *CssS*, *trpA*, *trpB*, *trpC*, *trpD*, *trpE*, *trpF*, *tdh/kbl*, *alsD*, *sigD*, *prpC*, *gapB*, ***pckA***, *fbp*, *rocA*, *ycgN*, *ycgM*, *rocF*, and *rocD*; and b) growing said altered *Bacillus* strain under conditions such that said protein of interest is expressed by said altered *Bacillus* strain, wherein said expression of said protein of interest is enhanced compared to the expression of said protein of interest in an unaltered *Bacillus* host strain, optionally wherein said protein of interest is a protease, optionally wherein a DNA construct is integrated into the chromosome of the *Bacillus* host cell under conditions such that said at least one gene is inactivated to produce an altered *Bacillus*.

Ferrari et al. further teach a method for obtaining an altered *Bacillus subtilis* strain with enhanced protease production comprising: a) transforming a *Bacillus subtilis* host cell with a DNA construct comprising at least one gene selected from the group consisting of *sbo*, *slr*, *ybcO*, *csn*, *spolISA*, *sigB*, *phrC*, *rapA*, *CssS*, *trpA*, *trpB*, *trpC*, *trpD*, *trpE*, *trpF*, *tdh/kbl*, *alsD*, *sigD*, *prpC*, *gapB*, *pckA*, *fbp*, *rocA*, *ycgN*, *ycgM*, *rocF*, and *rocD*,

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gene fragments thereof, and homologous sequences thereto, wherein said protein of interest in said DNA construct is a protease, and wherein said DNA construct is integrated into the chromosome of the *Bacillus subtilis* host cell under conditions such that said at least one gene is inactivated to produce an altered *Bacillus* strain; and b) growing said altered *Bacillus subtilis* strain under conditions such that enhanced protease production is obtained, optionally further comprising recovering said protease, which is selected from the group consisting of subtilisin 168, subtilisin BPN', subtilisin Carlsberg, subtilisin DY, subtilisin 147 and subtilisin 309 and variants thereof; optionally wherein said DNA construct comprises a pckA gene having the nucleic acid sequence of SEQ ID NO:27 encoding the amino acid sequence of SEQ ID NO: 28 (both sequences taught by Ferrari et al. are identical to Applicants' SEQ ID NOs: 27 and 28, see below alignment result and results in SCORE); optionally wherein said inactivation is by insertional inactivation of said at least one of said genes; optionally wherein said *Bacillus subtilis* strain further comprising at least one mutation in a gene selected from the group consisting of degU, degS, degQ, scoC4, spoIIIE, and oppA, wherein said mutation is degU(Hy)32; optionally wherein said altered *Bacillus subtilis* strain further comprises a deletion of one or more indigenous chromosomal regions or fragments thereof, wherein said indigenous chromosomal region includes about 0.5 to 500 kb. See Claims 1-88 of Ferrari et al.

Sequence alignment of Applicants' SEQ ID NO: 27 (Query) with SEQ ID NO: 27 taught by Ferrari et al. (Sbjct).

Score = 2920 bits (1581), Expect = 0.0
Identities = 1581/1581 (100%), Gaps = 0/1581 (0%)
Strand=Plus/Plus

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Query	1	ATGAACTCAGTTGATTTGACCGCTGATTTACAAGCCTTATTAACATGTCCAAATGTGCGT	60
Sbjct	1	ATGAACTCAGTTGATTTGACCGCTGATTTACAAGCCTTATTAACATGTCCAAATGTGCGT	60
Query	61	CATAATTTATCAGCAGCACAGCTAACAGAAAAAGTCCTCTCCCGAAACGAAGGCATTTTA	
120			
Sbjct	61	CATAATTTATCAGCAGCACAGCTAACAGAAAAAGTCCTCTCCCGAAACGAAGGCATTTTA	
120			
Query	121	ACATCCACAGGTGCTGTTTCGCGCGACAACAGGCGCTTACACAGGACGCTCACCTAAAGAT	
180			
Sbjct	121	ACATCCACAGGTGCTGTTTCGCGCGACAACAGGCGCTTACACAGGACGCTCACCTAAAGAT	
180			
Query	181	AAATTCATCGTGGAGGAAGAAAGCACGAAAAATAAGATCGATTGGGGCCCGGTGAATCAG	
240			
Sbjct	181	AAATTCATCGTGGAGGAAGAAAGCACGAAAAATAAGATCGATTGGGGCCCGGTGAATCAG	
240			
Query	241	CCGATTTTCAAGAAGCGTTTGAGCGGCTGTACACGAAAGTTGTCAGCTATTTAAAGGAG	
300			
Sbjct	241	CCGATTTTCAAGAAGCGTTTGAGCGGCTGTACACGAAAGTTGTCAGCTATTTAAAGGAG	
300			
Query	301	CGAGATGAACTGTTTGTTCGAAGGATTTGCCGGAGCAGACGAGAAATACAGGCTGCCG	
360			
Sbjct	301	CGAGATGAACTGTTTGTTCGAAGGATTTGCCGGAGCAGACGAGAAATACAGGCTGCCG	
360			
Query	361	ATCACTGTCGTAAATGAGTTCGCATGGCACAATTTATTTGCGCGGCAGCTGTTTATCCGT	
420			
Sbjct	361	ATCACTGTCGTAAATGAGTTCGCATGGCACAATTTATTTGCGCGGCAGCTGTTTATCCGT	
420			
Query	421	CCGGAAGGAAATGATAAGAAAACAGTTGAGCAGCCGTTACCATTCCTTTCTGCTCCGCAT	
480			
Sbjct	421	CCGGAAGGAAATGATAAGAAAACAGTTGAGCAGCCGTTACCATTCCTTTCTGCTCCGCAT	
480			
Query	481	TTCAAAGCGGATCCAAAAACAGACGGCACTCATTCCGAAACGTTTATTATTGTCTCTTTC	
540			
Sbjct	481	TTCAAAGCGGATCCAAAAACAGACGGCACTCATTCCGAAACGTTTATTATTGTCTCTTTC	
540			
Query	541	GAAAAGCGGACAATTTTAATCGGCGGAACTGAGTATGCCGGTGAAATGAAGAAGTCCATT	
600			
Sbjct	541	GAAAAGCGGACAATTTTAATCGGCGGAACTGAGTATGCCGGTGAAATGAAGAAGTCCATT	
600			
Query	601	TTCTCCATTATGAATTTCTTGCTGCCTGAAAGAGATATTTTATCTATGCACTGCTCCGCC	
660			
Sbjct	601	TTCTCCATTATGAATTTCTTGCTGCCTGAAAGAGATATTTTATCTATGCACTGCTCCGCC	
660			

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Query	661	AATGTCGGTGAAAAAGGCGATGTCGCCCTTTTCTTCGGACTGTCAGGAACAGGAAAGACC
720		
Sbjct	661	AATGTCGGTGAAAAAGGCGATGTCGCCCTTTTCTTCGGACTGTCAGGAACAGGAAAGACC
720		
Query	721	ACCCTGTCGGCAGATGCTGACCGCAAGCTGATCGGTGACGATGAACATGGCTGGTCTGAT
780		
Sbjct	721	ACCCTGTCGGCAGATGCTGACCGCAAGCTGATCGGTGACGATGAACATGGCTGGTCTGAT
780		
Query	781	ACAGGCGTCTTTAATATTGAAGGCGGATGCTACGCTAAGTGTATTTCATTTAAGCGAGGAA
840		
Sbjct	781	ACAGGCGTCTTTAATATTGAAGGCGGATGCTACGCTAAGTGTATTTCATTTAAGCGAGGAA
840		
Query	841	AAGGAGCCGCAAATCTTTAACGCGATCCGCTTCGGGTCTGTTCTCGAAAATGTCGTTGTG
900		
Sbjct	841	AAGGAGCCGCAAATCTTTAACGCGATCCGCTTCGGGTCTGTTCTCGAAAATGTCGTTGTG
900		
Query	901	GATGAAGATACACGCGAAGCCAATTATGATGATTCCTTCTATACTGAAAACACGCGGGCA
960		
Sbjct	901	GATGAAGATACACGCGAAGCCAATTATGATGATTCCTTCTATACTGAAAACACGCGGGCA
960		
Query	961	GCTTACCCGATTTCATATGATTAATAACATCGTGACTCCAAGCATGGCCGGCCATCCGTCA
1020		
Sbjct	961	GCTTACCCGATTTCATATGATTAATAACATCGTGACTCCAAGCATGGCCGGCCATCCGTCA
1020		
Query	1021	GCCATTGTATTTTTGACGGCTGATGCCTTCGGAGTCCTGCCGCCGATCAGCAAACCTAACG
1080		
Sbjct	1021	GCCATTGTATTTTTGACGGCTGATGCCTTCGGAGTCCTGCCGCCGATCAGCAAACCTAACG
1080		
Query	1081	AAGGAGCAGGTGATGTACCATTTTTTGAGCGGTTACACGAGTAAGCTTGCCGGAACCGAA
1140		
Sbjct	1081	AAGGAGCAGGTGATGTACCATTTTTTGAGCGGTTACACGAGTAAGCTTGCCGGAACCGAA
1140		
Query	1141	CGTGGTGTACGTCTCCTGAAACGACGTTTTCTACATGCTTCGGCTCACCGTTCCTGCCG
1200		
Sbjct	1141	CGTGGTGTACGTCTCCTGAAACGACGTTTTCTACATGCTTCGGCTCACCGTTCCTGCCG
1200		
Query	1201	CTTCCTGCTCACGTCTATGCTGAAATGCTCGGCAAAAAGATCGATGAACACGGCGCAGAC
1260		
Sbjct	1201	CTTCCTGCTCACGTCTATGCTGAAATGCTCGGCAAAAAGATCGATGAACACGGCGCAGAC
1260		
Query	1261	GTTTTCTTAGTCAATACCGGATGGACCGGGGGCGGCTACGGCACAGGCGAACGAATGAAG
1320		

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Sbjct  1261  GTTTTCTTAGTCAATACCGGATGGACCGGGGGCGGCTACGGCACAGGCGAACGAATGAAG
1320

Query  1321  CTTTCTTACACTAGAGCAATGGTCAAAGCAGCGATTGAAGGCAAATTAGAGGATGCTGAA
1380  ||||||||||||||||||||||||||||||||||||||||||||||||||||||||||||
Sbjct  1321  CTTTCTTACACTAGAGCAATGGTCAAAGCAGCGATTGAAGGCAAATTAGAGGATGCTGAA
1380

Query  1381  ATGATAACTGACGATATTTTCGGCCTGCACATTCCGGCCCATGTTCTGGCGTTCCTGAT
1440  ||||||||||||||||||||||||||||||||||||||||||||||||||||||||||||
Sbjct  1381  ATGATAACTGACGATATTTTCGGCCTGCACATTCCGGCCCATGTTCTGGCGTTCCTGAT
1440

Query  1441  CATATCCTTCAGCCTGAAAACACGTGGACCAACAAGGAAGAATACAAAGAAAAAGCAGTC
1500  ||||||||||||||||||||||||||||||||||||||||||||||||||||||||||||
Sbjct  1441  CATATCCTTCAGCCTGAAAACACGTGGACCAACAAGGAAGAATACAAAGAAAAAGCAGTC
1500

Query  1501  TACCTTGCAAATGAATTCAAAGAGAACTTTAAAAAGTTTCGCACATACCGATGCCATCGCC
1560  ||||||||||||||||||||||||||||||||||||||||||||||||||||||||||||
Sbjct  1501  TACCTTGCAAATGAATTCAAAGAGAACTTTAAAAAGTTTCGCACATACCGATGCCATCGCC
1560

Query  1561  CAGGCAGGCGGCCCTCTCGTA  1581
      ||||||||||||||||||||
Sbjct  1561  CAGGCAGGCGGCCCTCTCGTA  1581

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Sequence alignment of Applicants' SEQ ID NO: 28 (Query) with SEQ ID NO: 28

taught by Ferrari et al. (Sbjct).

Score = 1093 bits (2828), Expect = 0.0, Method: Compositional matrix adjust.
 Identities = 527/527 (100%), Positives = 527/527 (100%), Gaps = 0/527 (0%)

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Query  1      MNSVDLTADLQALLTCPNVRHNL SAAQLTEKVLSRNEGILTSTGAVRATTGAYTGRSPKD  60
Sbjct  1      MNSVDLTADLQALLTCPNVRHNL SAAQLTEKVLSRNEGILTSTGAVRATTGAYTGRSPKD  60

Query  61      KFIVEEESTKNKIDWGPVNQPISEEA FERLYTKVVSYLKERDEL FVFEGFAGADEKYRLP  120
Sbjct  61      KFIVEEESTKNKIDWGPVNQPISEEA FERLYTKVVSYLKERDEL FVFEGFAGADEKYRLP  120

Query  121     ITVVNEFAWHNLFARQLFIRPEGNDKKTVEQPFTILSAPHFKADPKTDGTHSETFIIVSF  180
Sbjct  121     ITVVNEFAWHNLFARQLFIRPEGNDKKTVEQPFTILSAPHFKADPKTDGTHSETFIIVSF  180

Query  181     EKRTILIGGTEYAGEMKKSIFSIMN FLLPERDILSMHCSANVGEKGDVALFFGLSGTGKT  240
Sbjct  181     EKRTILIGGTEYAGEMKKSIFSIMN FLLPERDILSMHCSANVGEKGDVALFFGLSGTGKT  240

Query  241     TLSADADRKLIGDDEHGWSDTGVFNIEGGCYAKCIHLSEEKEPQIFNAIRFGSVLENVVV  300
Sbjct  241     TLSADADRKLIGDDEHGWSDTGVFNIEGGCYAKCIHLSEEKEPQIFNAIRFGSVLENVVV

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Sbjct	241	TL SADADRKLIGDDEHGWSDTGVFNIEGGCYAKCIHLSEEKEPQIFNAIRFGSVLENVVV	300
Query	301	DEDTREANYDDSFYTENTRAAYPIHMINNIVTPSMAGHPSAIVFLTADAFGVLPPI SKLT	360
		DEDTREANYDDSFYTENTRAAYPIHMINNIVTPSMAGHPSAIVFLTADAFGVLPPI SKLT	
Sbjct	301	DEDTREANYDDSFYTENTRAAYPIHMINNIVTPSMAGHPSAIVFLTADAFGVLPPI SKLT	360
Query	361	KEQVMYHFLSGYTSKLAGTERGVTS PETTFSTCFGSPFLPLPAHVYAEMLGKKIDEHGAD	420
		KEQVMYHFLSGYTSKLAGTERGVTS PETTFSTCFGSPFLPLPAHVYAEMLGKKIDEHGAD	
Sbjct	361	KEQVMYHFLSGYTSKLAGTERGVTS PETTFSTCFGSPFLPLPAHVYAEMLGKKIDEHGAD	420
Query	421	VFLVNTGWTGGGYGTGERMKLSYTRAMV KAAIEGKLEDAEMITDDIFGLHIPAHVPGVPD	480
		VFLVNTGWTGGGYGTGERMKLSYTRAMV KAAIEGKLEDAEMITDDIFGLHIPAHVPGVPD	
Sbjct	421	VFLVNTGWTGGGYGTGERMKLSYTRAMV KAAIEGKLEDAEMITDDIFGLHIPAHVPGVPD	480
Query	481	HILQPENTWTNKEEYKEKAVYLANEFKENFKKFAHTDAIAQAGGPLV	527
		HILQPENTWTNKEEYKEKAVYLANEFKENFKKFAHTDAIAQAGGPLV	
Sbjct	481	HILQPENTWTNKEEYKEKAVYLANEFKENFKKFAHTDAIAQAGGPLV	527

Therefore, the teachings of Ferrari et al. anticipate Claims 38-44 and 55-64 for the reasons provided herein.

Claims 38-44 and 55-64 are rejected under 35 U.S.C. § 102(e) as being anticipated by Ferrari et al. (US Patent Application Publication No. US 2006/0073559 A1 filed on 03/28/2003).

The instant claims are drawn to [1] a method for enhancing production of at least one protein by a microorganism, comprising the steps: a) providing a microorganism host cell; b) inactivating the *pckA* gene in said host cell to produce an altered strain; and c) growing said altered strain under growth conditions suitable for expression of said protein; and [2] a method for obtaining *Bacillus subtilis* strains having enhanced protease production comprising the steps of: transforming a *B. subtilis* host cell with a DNA construct; allowing homologous recombination of the DNA construct and a homologous region of the *B. subtilis* chromosome wherein *pckA* is deleted from the *B.*

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subtilis chromosome to produce an altered *Bacillus subtilis* strain; and growing the altered *B. subtilis* strain under conditions suitable for the expression of said protease.

It is noted by the Examiner that the term, “homologous proteins” and “heterologous proteins” are defined in page 38 of the specification, and have been interpreted by the Examiner as how they are defined in the specification.

Ferrari et al. teach a method for enhancing expression of a protein of interest from *Bacillus* comprising: a) obtaining an altered *Bacillus* strain capable of producing a protein of interest, wherein said altered *Bacillus* strain has at least one inactivated chromosomal gene selected from the group consisting of *sbo*, *sir*, *ybcO*, *csn*, *spolISA*, *sigB*, *phrC*, *rapA*, *CssS*, *trpA*, *trpB*, *trpC*, *trpD*, *trpE*, *trpF*, *tdh/kbl*, *alsD*, *sigD*, *prpC*, *gapB*, ***pckA***, *fbp*, *rocA*, *ycgN*, *ycgM*, *rocF*, and *rocD*; and b) growing said altered *Bacillus* strain under conditions such that said protein of interest is expressed by said altered *Bacillus* strain, wherein said expression of said protein of interest is enhanced compared to the expression of said protein of interest in an unaltered *Bacillus* host strain, optionally wherein said protein of interest is a protease, optionally wherein a DNA construct is integrated into the chromosome of the *Bacillus* host cell under conditions such that said at least one gene is inactivated to produce an altered *Bacillus*.

Ferrari et al. further teach a method for obtaining an altered *Bacillus subtilis* strain with enhanced protease production comprising: a) transforming a *Bacillus subtilis* host cell with a DNA construct comprising at least one gene selected from the group consisting of *sbo*, *slr*, *ybcO*, *csn*, *spolISA*, *sigB*, *phrC*, *rapA*, *CssS*, *trpA*, *trpB*, *trpC*, *trpD*, *trpE*, *trpF*, *tdh/kbl*, *alsD*, *sigD*, *prpC*, *gapB*, *pckA*, *fbp*, *rocA*, *ycgN*, *ycgM*, *rocF*, and *rocD*,

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gene fragments thereof, and homologous sequences thereto, wherein said protein of interest in said DNA construct is a protease, and wherein said DNA construct is integrated into the chromosome of the *Bacillus subtilis* host cell under conditions such that said at least one gene is inactivated to produce an altered *Bacillus* strain; and b) growing said altered *Bacillus subtilis* strain under conditions such that enhanced protease production is obtained, optionally further comprising recovering said protease, which is selected from the group consisting of subtilisin 168, subtilisin BPN', subtilisin Carlsberg, subtilisin DY, subtilisin 147 and subtilisin 309 and variants thereof; optionally wherein said DNA construct comprises a pckA gene having the nucleic acid sequence of SEQ ID NO:27 encoding the amino acid sequence of SEQ ID NO: 28 (both sequences taught by Ferrari et al. are identical to Applicants' SEQ ID NOs: 27 and 28, see below alignment result and results in SCORE); optionally wherein said inactivation is by insertional inactivation of said at least one of said genes; optionally wherein said *Bacillus subtilis* strain further comprising at least one mutation in a gene selected from the group consisting of degU, degS, degQ, scoC4, spoIIE, and oppA, wherein said mutation is degU(Hy)32; optionally wherein said altered *Bacillus subtilis* strain further comprises a deletion of one or more indigenous chromosomal regions or fragments thereof, wherein said indigenous chromosomal region includes about 0.5 to 500 kb. See Claims 1-88 of Ferrari et al.

Sequence alignment of Applicants' SEQ ID NO: 27 (Query) with SEQ ID NO: 27 taught by Ferrari et al. (Sbjct).

Score = 2920 bits (1581), Expect = 0.0
Identities = 1581/1581 (100%), Gaps = 0/1581 (0%)
Strand=Plus/Plus

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Query	1	ATGAACTCAGTTGATTTGACCGCTGATTTACAAGCCTTATTAACATGTCCAAATGTGCGT	60
Sbjct	1	ATGAACTCAGTTGATTTGACCGCTGATTTACAAGCCTTATTAACATGTCCAAATGTGCGT	60
Query	61	CATAATTTATCAGCAGCACAGCTAACAGAAAAAGTCCTCTCCCGAAACGAAGGCATTTTA	
120			
Sbjct	61	CATAATTTATCAGCAGCACAGCTAACAGAAAAAGTCCTCTCCCGAAACGAAGGCATTTTA	
120			
Query	121	ACATCCACAGGTGCTGTTTCGCGCGACAACAGGCGCTTACACAGGACGCTCACCTAAAGAT	
180			
Sbjct	121	ACATCCACAGGTGCTGTTTCGCGCGACAACAGGCGCTTACACAGGACGCTCACCTAAAGAT	
180			
Query	181	AAATTCATCGTGGAGGAAGAAAGCACGAAAAATAAGATCGATTGGGGCCCGGTGAATCAG	
240			
Sbjct	181	AAATTCATCGTGGAGGAAGAAAGCACGAAAAATAAGATCGATTGGGGCCCGGTGAATCAG	
240			
Query	241	CCGATTTTCAGAAGAAGCGTTTGAGCGGCTGTACACGAAAGTTGTCAGCTATTTAAAGGAG	
300			
Sbjct	241	CCGATTTTCAGAAGAAGCGTTTGAGCGGCTGTACACGAAAGTTGTCAGCTATTTAAAGGAG	
300			
Query	301	CGAGATGAACTGTTTGTTCGAAGGATTTGCCGGAGCAGACGAGAAATACAGGCTGCCG	
360			
Sbjct	301	CGAGATGAACTGTTTGTTCGAAGGATTTGCCGGAGCAGACGAGAAATACAGGCTGCCG	
360			
Query	361	ATCACTGTCGTAAATGAGTTCGCATGGCACAATTTATTTGCGCGGCAGCTGTTTATCCGT	
420			
Sbjct	361	ATCACTGTCGTAAATGAGTTCGCATGGCACAATTTATTTGCGCGGCAGCTGTTTATCCGT	
420			
Query	421	CCGGAAGGAAATGATAAGAAAACAGTTGAGCAGCCGTTACCATTCCTTTCTGCTCCGCAT	
480			
Sbjct	421	CCGGAAGGAAATGATAAGAAAACAGTTGAGCAGCCGTTACCATTCCTTTCTGCTCCGCAT	
480			
Query	481	TTCAAAGCGGATCCAAAAACAGACGGCACTCATTCCGAAACGTTTATTATTGTCTCTTTC	
540			
Sbjct	481	TTCAAAGCGGATCCAAAAACAGACGGCACTCATTCCGAAACGTTTATTATTGTCTCTTTC	
540			
Query	541	GAAAAGCGGACAATTTTAATCGGCGGAACTGAGTATGCCGGTGAAATGAAGAAGTCCATT	
600			
Sbjct	541	GAAAAGCGGACAATTTTAATCGGCGGAACTGAGTATGCCGGTGAAATGAAGAAGTCCATT	
600			
Query	601	TTCTCCATTATGAATTTCTTGCTGCCTGAAAGAGATATTTTATCTATGCACTGCTCCGCC	
660			
Sbjct	601	TTCTCCATTATGAATTTCTTGCTGCCTGAAAGAGATATTTTATCTATGCACTGCTCCGCC	
660			

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Query 661 AATGTCGGTGAAAAAGGCGATGTCGCCCTTTTCTTCGGACTGTCAGGAACAGGAAAGACC
720 |||||
Sbjct 661 AATGTCGGTGAAAAAGGCGATGTCGCCCTTTTCTTCGGACTGTCAGGAACAGGAAAGACC
720

Query 721 ACCCTGTCGGCAGATGCTGACCGCAAGCTGATCGGTGACGATGAACATGGCTGGTCTGAT
780 |||||
Sbjct 721 ACCCTGTCGGCAGATGCTGACCGCAAGCTGATCGGTGACGATGAACATGGCTGGTCTGAT
780

Query 781 ACAGGCGTCTTTAATATTGAAGGCGGATGCTACGCTAAGTGTATTTCATTTAAGCGAGGAA
840 |||||
Sbjct 781 ACAGGCGTCTTTAATATTGAAGGCGGATGCTACGCTAAGTGTATTTCATTTAAGCGAGGAA
840

Query 841 AAGGAGCCGCAAATCTTTAACGCGATCCGCTTCGGGTCTGTTCTCGAAAATGTCGTTGTG
900 |||||
Sbjct 841 AAGGAGCCGCAAATCTTTAACGCGATCCGCTTCGGGTCTGTTCTCGAAAATGTCGTTGTG
900

Query 901 GATGAAGATACACGCGAAGCCAATTATGATGATTCCTTCTATACTGAAAACACGCGGGCA
960 |||||
Sbjct 901 GATGAAGATACACGCGAAGCCAATTATGATGATTCCTTCTATACTGAAAACACGCGGGCA
960

Query 961 GCTTACCCGATTTCATATGATTAATAACATCGTGACTCCAAGCATGGCCGGCCATCCGTCA
1020 |||||
Sbjct 961 GCTTACCCGATTTCATATGATTAATAACATCGTGACTCCAAGCATGGCCGGCCATCCGTCA
1020

Query 1021 GCCATTGTATTTTTGACGGCTGATGCCTTCGGAGTCCTGCCGCCGATCAGCAAACCTAACG
1080 |||||
Sbjct 1021 GCCATTGTATTTTTGACGGCTGATGCCTTCGGAGTCCTGCCGCCGATCAGCAAACCTAACG
1080

Query 1081 AAGGAGCAGGTGATGTACCATTTTTTGAGCGGTTACACGAGTAAGCTTGCCGGAACCGAA
1140 |||||
Sbjct 1081 AAGGAGCAGGTGATGTACCATTTTTTGAGCGGTTACACGAGTAAGCTTGCCGGAACCGAA
1140

Query 1141 CGTGGTGTACGTCTCCTGAAACGACGTTTTCTACATGCTTCGGCTCACCGTTCCTGCCG
1200 |||||
Sbjct 1141 CGTGGTGTACGTCTCCTGAAACGACGTTTTCTACATGCTTCGGCTCACCGTTCCTGCCG
1200

Query 1201 CTTCTGCTCACGTCTATGCTGAAATGCTCGGCAAAAAGATCGATGAACACGGCGCAGAC
1260 |||||
Sbjct 1201 CTTCTGCTCACGTCTATGCTGAAATGCTCGGCAAAAAGATCGATGAACACGGCGCAGAC
1260

Query 1261 GTTTTCTTAGTCAATACCGGATGGACCGGGGGCGGCTACGGCACAGGCGAACGAATGAAG
1320 |||||

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Sbjct  1261  GTTTTCTTAGTCAATACCGGATGGACCGGGGGCGGCTACGGCACAGGCGAACGAATGAAG
1320

Query  1321  CTTTCTTACACTAGAGCAATGGTCAAAGCAGCGATTGAAGGCAAATTAGAGGATGCTGAA
1380  ||||||||||||||||||||||||||||||||||||||||||||||||||||||||||||
Sbjct  1321  CTTTCTTACACTAGAGCAATGGTCAAAGCAGCGATTGAAGGCAAATTAGAGGATGCTGAA
1380

Query  1381  ATGATAACTGACGATATTTTCGGCCTGCACATTCCGGCCCATGTTCTGGCGTTCCTGAT
1440  ||||||||||||||||||||||||||||||||||||||||||||||||||||||||||||
Sbjct  1381  ATGATAACTGACGATATTTTCGGCCTGCACATTCCGGCCCATGTTCTGGCGTTCCTGAT
1440

Query  1441  CATATCCTTCAGCCTGAAAACACGTGGACCAACAAGGAAGAATACAAAGAAAAAGCAGTC
1500  ||||||||||||||||||||||||||||||||||||||||||||||||||||||||||||
Sbjct  1441  CATATCCTTCAGCCTGAAAACACGTGGACCAACAAGGAAGAATACAAAGAAAAAGCAGTC
1500

Query  1501  TACCTTGCAAATGAATTCAAAGAGAACTTTAAAAAGTTTCGCACATACCGATGCCATCGCC
1560  ||||||||||||||||||||||||||||||||||||||||||||||||||||||||||||
Sbjct  1501  TACCTTGCAAATGAATTCAAAGAGAACTTTAAAAAGTTTCGCACATACCGATGCCATCGCC
1560

Query  1561  CAGGCAGGCGGCCCTCTCGTA  1581
      ||||||||||||||||||||
Sbjct  1561  CAGGCAGGCGGCCCTCTCGTA  1581

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Sequence alignment of Applicants' SEQ ID NO: 28 (Query) with SEQ ID NO: 28
taught by Ferrari et al. (Sbjct).

Score = 1093 bits (2828), Expect = 0.0, Method: Compositional matrix adjust.
Identities = 527/527 (100%), Positives = 527/527 (100%), Gaps = 0/527 (0%)

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Query  1      MNSVDLTADLQALLTCPNVRHNLSAAQLTEKVLSRNEGILTSTGAVRATTGAYTGRSPKD  60
Sbjct  1      MNSVDLTADLQALLTCPNVRHNLSAAQLTEKVLSRNEGILTSTGAVRATTGAYTGRSPKD  60

Query  61      KFIVEEESTKNKIDWGPVNQPISEEAFERLYTKVVSYLKERDELFFVEGFAGADEKYRLP  120
Sbjct  61      KFIVEEESTKNKIDWGPVNQPISEEAFERLYTKVVSYLKERDELFFVEGFAGADEKYRLP  120

Query  121     ITVVNEFAWHNLFARQLFIRPEGNDKKTVEQPFTILSAPHFKADPKTDGTHSETFIIVSF  180
Sbjct  121     ITVVNEFAWHNLFARQLFIRPEGNDKKTVEQPFTILSAPHFKADPKTDGTHSETFIIVSF  180

Query  181     EKRTILIGGTEYAGEMKKSIFSIMNPLLPERDILSMHCSANVGEKGDVALFFGLSGTGKT  240
Sbjct  181     EKRTILIGGTEYAGEMKKSIFSIMNPLLPERDILSMHCSANVGEKGDVALFFGLSGTGKT  240

Query  241     TLSADADRKLIGDDEHGWSDTGVFNIEGGCYAKCIHLSEEKEPQIFNAIRFGSVLENNVV  300
Sbjct  241     TLSADADRKLIGDDEHGWSDTGVFNIEGGCYAKCIHLSEEKEPQIFNAIRFGSVLENNVV

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Sbjct	241	TLSADADRKLIGDDEHGWSDTGVFNIEGGCYAKCIHLSEEKEPQIFNAIRFGSVLENVVV	300
Query	301	DEDTREANYDDSFYTENTRAAYPIHMINNIVTPSMAGHPSAIVFLTADAFGVLPPI SKLT	360
		DEDTREANYDDSFYTENTRAAYPIHMINNIVTPSMAGHPSAIVFLTADAFGVLPPI SKLT	
Sbjct	301	DEDTREANYDDSFYTENTRAAYPIHMINNIVTPSMAGHPSAIVFLTADAFGVLPPI SKLT	360
Query	361	KEQVMYHFLSGYTSKLAGTERGVTS PETTFSTCFGSPFLPLPAHVYAEMLGKKIDEHGAD	420
		KEQVMYHFLSGYTSKLAGTERGVTS PETTFSTCFGSPFLPLPAHVYAEMLGKKIDEHGAD	
Sbjct	361	KEQVMYHFLSGYTSKLAGTERGVTS PETTFSTCFGSPFLPLPAHVYAEMLGKKIDEHGAD	420
Query	421	VFLVNTGWTGGGYGTGERMKLSYTRAMVKAAIEGKLEDAEMITDDIFGLHIPAHVPGVPD	480
		VFLVNTGWTGGGYGTGERMKLSYTRAMVKAAIEGKLEDAEMITDDIFGLHIPAHVPGVPD	
Sbjct	421	VFLVNTGWTGGGYGTGERMKLSYTRAMVKAAIEGKLEDAEMITDDIFGLHIPAHVPGVPD	480
Query	481	HILQPENTWTNKEEYKEKAVYLANEFKENFKKFAHTDAIAQAGGPLV	527
		HILQPENTWTNKEEYKEKAVYLANEFKENFKKFAHTDAIAQAGGPLV	
Sbjct	481	HILQPENTWTNKEEYKEKAVYLANEFKENFKKFAHTDAIAQAGGPLV	527

Therefore, the teachings of Ferrari et al. anticipate Claims 38-44 and 55-64 for the reasons provided herein.

Conclusion

Claims 38-44 and 55-64 are rejected for the reasons as stated above. Applicants must respond to the objections/rejections in this Office action to be fully responsive in prosecution.

The instant Office action is non-final.

Any inquiry concerning this communication or earlier communications from the examiner should be directed to Jae W. Lee whose telephone number is 571-272-9949. The examiner can normally be reached on M-F between 9:00-6:00.

If attempts to reach the examiner by telephone are unsuccessful, the examiner's supervisor, Manjunath Rao can be reached on 571-272-0939. The fax phone number for the organization where this application or proceeding is assigned is 571-273-8300.

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/JAE W LEE/
Examiner, Art Unit 1656

/SUZANNE M NOAKES/
Primary Examiner, Art Unit 1656